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The Pfizer Revelation: A Call for Accountability and Immediate Global Withdrawal

The launch of the Pfizer-BioNTech mRNA vaccine in December 2020 was framed as a milestone in public health, yet internal documents declassified in October 2021 tell a significantly different story. According to the sources, a confidential report titled “Cumulative Analysis of Post-Authorization Adverse Event Reports” reveals that within the first 90 days of the rollout, Pfizer had already documented a staggering wave of mortality and morbidity. Despite this internal data, which recorded 1,223 deaths and 158,893 adverse events by February 28, 2021, global health agencies continued to endorse the vaccine as safe and effective.

The data, which the sources describe as a “de facto Mea Culpa” from Pfizer, provides irrefutable evidence of the injection’s risks because it emanates directly from the manufacturer’s own records. Among the most alarming findings were 23 spontaneous abortions out of 270 pregnancies and over 2,000 reports of cardiac disorders. Specific cardiovascular evaluations in the report identified 1,403 cases of serious heart-related issues, with a median onset of symptoms occurring less than 24 hours after injection. Furthermore, the analysis of these sources suggests that the FDA, CDC, and NIAID were fully aware of these dangers as early as March 2021 but chose to withhold this information from the public, even attempting to keep the data hidden for 55 years.

The sources highlight a profound institutional failure, noting that it took until August 2025 for the FDA to finally order Pfizer and Moderna to issue warnings regarding the “high risk of heart damage” and myocarditis, particularly in young men. This warning came more than four years after the risk was documented in the internal 2021 report. This delay is characterized in the sources as a betrayal of public trust and a “criminal undertaking” by both Big Pharma and complicit government entities. To provide context for these actions, the sources point to Pfizer’s history, specifically a $2.3 billion medical fraud settlement in 2009 involving charges of fraudulent marketing.

In light of these revelations, the sources present a persuasive case for the immediate worldwide withdrawal of the mRNA vaccine. The narrative asserts that the vaccine was never intended to save lives but has instead functioned as a multibillion-dollar operation that persists despite documented evidence of harm. The sources conclude by calling upon the general public and professionals to perusal the declassified 38-page report and its appendices to verify these claims, urging a global cancellation of the vaccination program to prevent further injury and loss of life.

Analogy for Understanding: To grasp the severity of these findings, consider an airline that discovers a fatal structural flaw in a new aircraft during its first three months of flight. If the airline and the aviation regulators chose to hide those crash reports while mandating that every citizen must fly on that specific plane, the eventual "declassification" of those safety logs would be viewed as an undeniable admission of negligence and a total collapse of safety standards. This report is presented as the safety log for the mRNA project.